



# **Bombshell: Pfizer's "Secret" 2021 Declassified Report. CALL FOR IMMEDIATE WITHDRAWAL OF COVID "VACCINE"**

## **Introduction: The Launch of the mRNA Project and Initial Institutional Awareness**

The Pfizer BioNTech mRNA vaccine was officially launched in the United States on December 14, 2020, following the granting of an Emergency Use Authorization (EUA) on December 11, 2020. According to documents and analysis provided by Professor Michel Chossudovsky, several key health agencies within the U.S. Department of Health and Human Services (HHS) were aware of the potential underlying dangers of this vaccine from its inception. These agencies included the Center for Disease Control (CDC), the Food and Drug Administration (FDA), and the National Institute of Allergy and Infectious Diseases (NIAID), then led by Dr. Anthony Fauci.

Despite this internal awareness, these organizations, in coordination with the World Health Organization (WHO) and under reported pressure from the pharmaceutical industry, granted the "greenlight" for Pfizer and Moderna to market mRNA vaccines on a global scale. To monitor the early impacts of the rollout, Pfizer commissioned a detailed, originally classified report entitled "Cumulative Analysis of Post-Authorization Adverse Event Reports of PF-07302048 (BNT162B2) Received Through 28-Feb-2021". The primary objective of this report was to tabulate and categorize data regarding the vaccine's impact on mortality, morbidity, and various adverse events during the initial two and a half months of its distribution, spanning from mid-December 2020 to the end of February 2021.

## **The Declassification of the Pfizer Report**

The findings contained within this classified report were not initially disclosed to the general public. Instead, they were shared with the specific health agencies responsible for overseeing the vaccine distribution under the EUA. It was not until October 2021 that this document was declassified following a Freedom of Information (FOI) procedure. The release of this data has been described as a "bombshell" because it revealed that by late February 2021, Pfizer had already recorded over 1,200 reports of deaths and thousands of adverse events allegedly linked to the vaccine.

The FDA reportedly sought to keep this data hidden from public view for a period of 55 years. The eventual "document dump" confirmed that within a very short reporting window of less than three months, the vaccine project was associated with significant safety concerns that contradicted the official narratives promoted by many governments and the WHO.

## **Comprehensive Statistical Analysis of Adverse Events**

The declassified data provides a stark overview of the vaccine's impact during the first 90 days of its global release. Between December 1, 2020, and February 28, 2021, Pfizer shipped an undisclosed number of doses worldwide. During this interval, the company received a cumulative total of 42,086 case reports. Within these reports, there were 158,893 individual adverse events documented.

The reports were categorized into two main types:

1. **Medically Confirmed Cases:** 25,379 reports.
2. **Non-Medically Confirmed Cases:** 16,707 reports.

The geographical distribution of these cases was widespread, though concentrated in a few key nations. The majority of cases (34,762) originated from:

- **United States:** 13,739 cases.
- **United Kingdom:** 13,404 cases.
- **Italy:** 2,578 cases.
- **Germany:** 1,913 cases.
- **France:** 1,506 cases.
- **Portugal:** 866 cases.
- **Spain:** 756 cases.

An additional 7,324 cases were distributed across 56 other countries. This data suggests that by early March 2021, health authorities were fully aware of an emerging wave of mortality and morbidity on a global scale.

## Demographics and Case Outcomes

The Pfizer report provided a detailed breakdown of the individuals affected by these adverse events, revealing significant trends in gender and age. According to Table 1 of the report, the gender distribution of reported cases was as follows:

- **Female:** 29,914 cases.
- **Male:** 9,182 cases.
- **Data Not Provided:** 2,990 cases.

The age range of those reporting events was vast, spanning from infants (0.01 years) to the elderly (107 years), with a mean age of 50.9 years. The specific age cohorts included:

- **17 Years and Younger:** 175 cases (including 46 cases under 16 and 34 cases under 12).
- **18 to 30 Years:** 4,953 cases.
- **31 to 50 Years:** 13,886 cases.
- **51 to 64 Years:** 7,884 cases.
- **65 to 74 Years:** 3,098 cases.
- **75 Years and Older:** 5,214 cases.
- **Unknown Age:** 6,876 cases.

The outcomes of these cases at the time of the report were equally concerning. While 19,582 individuals were listed as recovered or recovering, 11,361 had not recovered at the time the data was compiled. Most notably, the report documented **1,223 fatal outcomes** during this initial two-and-a-half-month period. Additionally, 520 individuals recovered but were left with sequelae (permanent or long-term conditions resulting from the event), and the status of 9,400 cases remained unknown.

## Classification of System Organ Class (SOC) Disorders

The adverse events were categorized by the physiological systems they impacted. The report highlighted several "System Organ Classes" that contained the highest number of events (greater than or equal to 2% of the total dataset).

**General Disorders and Administration Site Conditions** This was the most frequent category, accounting for 51,335 events. Specific symptoms included:

- **Pyrexia (Fever):** 7,666 cases (18.2%).
- **Fatigue:** 7,338 cases (17.4%).
- **Chills:** 5,514 cases (13.1%).

- **Vaccination Site Pain:** 5,181 cases (12.3%).

**Nervous System Disorders** There were 25,957 reported events related to the nervous system.

**Musculoskeletal and Connective Tissue Disorders** These accounted for 17,283 events.

**Gastrointestinal Disorders** A total of 14,096 gastrointestinal events were recorded. Specific symptoms included:

- **Nausea:** 5,182 cases (12.3%).
- **Diarrhea:** 1,880 cases (4.5%).
- **Vomiting:** 1,698 cases (4.0%).

#### Other Significant Categories

- **Respiratory, Thoracic, and Mediastinal Disorders:** 8,848 events.
- **Skin and Subcutaneous Tissue Disorders:** 8,476 events.
- **Injury, Poisoning, and Procedural Complications:** 5,590 events.
- **Infections and Infestations:** 4,610 events.
- **Investigations:** 3,693 events.

### Cardiovascular Adverse Events of Special Interest (AESIs)

Cardiovascular issues were a primary area of concern in the Pfizer report. The data revealed over **2,000 reports of cardiac disorders**. A more granular evaluation of cardiovascular AESIs (Table 7) identified 1,403 cases, which represented 3.3% of the total dataset. Of these, 241 were medically confirmed.

The specific cardiovascular conditions reported included:

- **Tachycardia:** 1,098 cases.
- **Arrhythmia:** 102 cases.
- **Myocardial Infarction (Heart Attack):** 89 cases.
- **Cardiac Failure:** 80 cases.
- **Acute Myocardial Infarction:** 41 cases.
- **Cardiac Failure Acute:** 11 cases.
- **Cardiogenic Shock and Postural Orthostatic Tachycardia Syndrome:** 7 cases each.
- **Coronary Artery Disease:** 6 cases.

The onset latency for these cardiovascular events was notably short, ranging from less than 24 hours to 21 days, with a median onset of less than 24 hours after injection.

### Impacts on Pregnancy and Spontaneous Abortions

The report also contained troubling data regarding pregnancy. Among the initial cohort of vaccinated individuals, there were 270 reported pregnancies. Out of these, the report documented **23 cases of spontaneous abortions**. This information has been highlighted as a critical data point that was available to health authorities early in the vaccination campaign but was not emphasized in public health communications.

### Institutional Response and the "Mea Culpa" Argument

The author of the analysis argues that the existence and subsequent declassification of this report constitute a "de facto Mea Culpa" on the part of Pfizer. Because the data emanates directly from the pharmaceutical company's own records, it is characterized as irrefutable evidence. The analysis suggests that because Pfizer was fully aware that

the mRNA vaccine would result in a significant wave of mortality and morbidity, the continued marketing of the product is tantamount to a "crime against humanity" and a "criminal undertaking".

Despite the "catastrophic nature" of the data revealed in the 2021 report, the author notes that as of early 2026, no meaningful attempts have been made by most national governments to withdraw the vaccine from the market. Furthermore, it is claimed that governments worldwide were pressured by Big Pharma to ignore these findings and maintain the official vaccine narrative.

## Historical Context: Pfizer's Record of Medical Fraud

To understand the current situation, the author points to Pfizer's historical record with the law. In 2009, Pfizer reached a \$2.3 billion settlement with the U.S. Department of Justice (DOJ) over charges of **"fraudulent marketing"**. As part of this settlement, Pfizer entered into an "expansive corporate integrity agreement" with the Office of Inspector General of the Department of Health and Human Services.

This agreement was intended to put procedures in place to detect and avoid future conduct similar to the fraud that led to the settlement. However, the author argues that the 2009 decision to place Pfizer on "probation" rather than pursuing more severe criminal penalties for its executives was a mistake, suggesting that the company's current actions regarding the mRNA vaccine are a continuation of past patterns of behavior.

## The 2025 FDA Warning and Retrospective Insights

In what is described as a "bitter irony," the analysis notes that by August 2025, the FDA finally ordered both Pfizer and Moderna to issue formal warnings regarding their mRNA vaccines. These new guidelines highlight a **high risk of heart damage**, specifically myocarditis, particularly among young men.

The author contends that this "deadly risk" was known to the FDA and the CDC from the very beginning, as evidenced by the 2021 classified report. The failure to act for several years, while instead endorsing and imposing the vaccine internationally, is presented as a major failure of public health institutions. The analysis asserts that the claim that the vaccine was intended to "save lives" was a falsehood, and that the operation has primarily served as a multibillion-dollar business for the pharmaceutical industry.

## Media Influence and Conflicts of Interest

The report also touches upon the role of the media in shaping public perception. It is claimed that mainstream "Fact Checkers" have ignored the declassified Pfizer report. The author highlights a specific potential conflict of interest involving James C. Smith, the former CEO of Reuters and Chairman of the Reuters "Fact Checker" board, who is also reportedly a top investor and board member of Pfizer. This connection is cited as a reason for the lack of critical media coverage regarding the adverse event data.

## Conclusion and Call for Action

The declassified Pfizer report, covering the period from December 2020 to February 2021, provides a detailed record of significant medical concerns that were documented almost immediately upon the vaccine's release. With 42,086 case reports and 1,223 deaths recorded in less than three months, the author calls for the **immediate withdrawal of the Covid mRNA vaccine**.

The data suggests that the physiological impacts of the vaccine—spanning from cardiac disorders and nervous system issues to spontaneous abortions—were documented and known to both the manufacturer and national health agencies early on. The author concludes that the ongoing distribution of the vaccine, despite this internal data, represents a global crisis and a fundamental violation of human safety and corporate integrity.

For those seeking to verify these claims, the author invites a thorough perusal of the 38-page declassified report and its various appendices, which contain the comprehensive tables and SOC classifications discussed herein. The final message of the analysis is one of urgent necessity: the cancellation of the vaccine program to prevent further mortality and morbidity.

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**Analogy for Understanding:** To understand the significance of this report, imagine a car manufacturer discovering during the first 90 days of a new model's release that the brakes were failing at an alarming rate, leading to hundreds of accidents. If that manufacturer and the safety regulators chose to keep those internal crash reports "classified" while continuing to tell the public the car was the safest ever made, the eventual "declassification" of those reports would be seen as the ultimate proof of negligence. This report is presented as that internal "crash data" for the vaccine project.